CLINICAL LABORATORY

SUSPECTED BIOTERRORISM (BT) EVENT MANAGEMENT GUIDELINE

Washington State Clinical Laboratory Advisory Council April 2003

Covert Event			Overt Event			
 LRN Sentinel Laboratory*: Unusual number of clinical section patients with similar section patients with similar section patients with similar section patients with similar section patients. Preliminary laboratory finding agent (see back page) Receipt of clinical specimen 	ymptoms gs suggestive of a E	Laboratory notified of increased level of suspicion	Notification from Public Health authorities, Infection Control, local media, etc. of potential bioterrorist threat			
BT agents • Unusual isolates from more t	han one patient		Customize the telephone numbers in the box below for YOUR laboratory			
Inform clinician of pertinent laboratory results and status of confirmatory testing	Inform Local Health Jurisdiction officials	Preserve and secure specimen/sample/all culture plates Wait for instructions from Local Health Jurisdiction for follow-up steps	Telephone Numbers for YOUR Facility: Laboratory Director Laboratory Supervisor Lead Technologist Infection Control Local Health Jurisdiction			

Local Health Jurisdictions will:

- > Inform and involve Washington State Department of Health (DOH) Epidemiology staff and the DOH Public Health Laboratories to determine where suspect samples are to be sent for further studies
- $> Advise\ LRN\ Sentinel\ laboratory\ on:\ -which\ LRN\ Reference **\ laboratory\ to\ send\ the\ specimen/sample$
 - -how to send the sample
 - -special packaging instructions
- * LRN Sentinel Laboratory: Laboratories that perform microbiology testing to RULE OUT a BT agent.
- ** LRN Reference Laboratory: Laboratories specifically authorized by the Centers for Disease Control and Prevention to perform testing to RULE IN the BT agent.

ENVIRONMENTAL SAMPLES: DO NOT ACCEPT any type of non-clinical specimen such as powders, other suspicious substances, or packages. Contact your Local Health Jurisdiction. REFER all phone calls from people regarding environmental specimens to local law enforcement or to your local health jurisdiction.

LRN SENTINEL LABORATORY **REFERENCE TABLE**

Agent	Culture Methods	Incubation Methods	Recovery Time	Colonial Morphology	Gram Stain Morphology	Preliminary Identification Tests	Action
Bacillus anthracis From: vesicle, sputum, CSF, blood, stool, rectal swab	Blood, Chocolate agar No growth on Mac	35°C in ambient air or CO ₂	8-24 hours	Non-hemolytic, gray colonies with ground glass appearance which "peaks" when touched	Large gram positive rods, oval, sub-terminal spores, no swelling of cell., capsules may be seen from specimen Gram stained	Catalase—positive Motility—negative	Refer to Laboratory designated by the local health jurisdiction
Francisella tularensis From: Blood, tissue, sputum, lymph nodes	Chocolate, BCYE, Thioglycollate, and Thayer-Martin agar Poor growth on BA No growth on Mac	35°C in CO ₂	~24-48 hours Hold up to 10 days	Very small, blue/gray colonies	Tiny gram negative coccobacilli poorly staining	Catalase—negative or weakly positive Oxidase—negative Urea—negative Motility—negative XV strip-no satelliting	Refer to Laboratory designated by the local health jurisdiction
Yersinia pestis From: Lymph node, blood, spleen, liver, sputum, bubo	Grows on routine culture media	22-28°C in ambient air or CO ₂	Grows slowly, 24-48 hours	Small, fried egg colonies may look like beaten copper	Gram negative rods which may show bi-polar staining	Catalase –positive Oxidase—negative Urea—negative Motility –negative TSI—weak acid slant, no change in butt	Refer to Laboratory designated by the local health jurisdiction
Brucella sp. From: Blood, bone marrow, tissue, CSF	Blood, Chocolate, Thayer-Martin or BCYE agar Some strains grow on Mac	35°C in CO ₂	Normally 24- 72 hours, may take up to 30 days	Small, gray/white colonies, punctate	Small gram negative cocco- bacilli, poorly staining	Catalase—positive Oxidase—positive Urease—positive XV—negative	Refer to Laboratory designated by the local health jurisdiction
Clostridium botulinum From: Feces, tissue, wound exudates, gastric contents	Blood or brucella agar Chopped meat Broth	Anaerobic incubation at 35°C	24-30 hours	Beta hemolytic with rhizoid colonies on moisture-free media; always swarms on damp media	Gram positive rods with oval, sub- terminal spores which swells the cell	Catalase—negative Indole—negative	Refer to Laboratory designated by the local health jurisdiction
Burkholderia pseudomallei & mallei From: Blood, sputum, wounds	Grows on routine culture media, strongly lactose + on Mac	35°C in ambient or CO ₂	24 hours B. mallei grows more slowly	Creamy tan to orange wrinkled colonies when old, fresh isolate may look like mercury	Gram negative rods similar to Pseudomonas	Catalase—positive Oxidase—positive B.mallei- var. oxidase/non- motile	Refer to Laboratory designated by the local health jurisdiction

- 1. Basic Diagnostic Testing Protocols for Level A Laboratories (updated: December 18, 2002). Centers for Disease Control and Prevention, American Society for Microbiology, and the Association of Public Health Laboratories.
- Biological and Chemical Terrorism: Strategic Plan for Preparedness and Response. CDC MMWR Volume 49/No.RR-4, April 21, 2001.
 Manual of Clinical Microbiology, 7th ed., American Society for Microbiology, 1999. Patrick R. Murray, editor-in-chief.
 USAMRIID's Medical Management of Biological Casualties, Handbook 4th ed. February, 2001 Appendix E.